

# Modelling the spatial organization of cell proliferation in the developing central nervous system

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## Abstract

How far is neuroepithelial cell proliferation in the developing central nervous system a deterministic process? Or, to put it in a more precise way, how accurately can it be described by a deterministic mathematical model? To provide tracks to answer this question, a deterministic system of transport and diffusion partial differential equations, both physiologically and spatially structured, is introduced as a model to describe the spatially organized process of cell proliferation during the development of the central nervous system. As an initial step towards dealing with the three-dimensional case, a unidimensional version of the model is presented. Numerical analysis and numerical tests are performed. In this work we also achieve a first experimental validation of the proposed model, by using cell proliferation data recorded from histological sections obtained during the development of the optic tectum in the chick embryo.

## 1 Introduction

The present work aims at modeling the spatial organization of the neuroepithelial (NE) cell proliferation in a developing cortical structure along an early and brief developmental period by using sets of quantitative data empirically obtained from a standardized experimental model: the developing chick optic tectum (OT). The mathematical model is based on a deterministic approach that uses the formalism of partial differential equations.

### 1.1 Biological background: Developmental neurobiology

#### 1.1.1 Relevance of a spatial and temporal organization in a developing system

The appropriate number of cells of each terminally differentiated cell type and also the spatial patterns they exhibit within the different tissues and organs composing

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pluricellular organisms are governed by interactive self-regulating behaviors that the developing cells exhibit during the embryonic development [1]. The increase in supra-cellular complexity generated during development requires the temporally and spatially organized operation of specific developmental cell behaviors (DCBs). These DCBs are usually reciprocally regulated and operate simultaneously and interactively [2]. Every developing cell population can be considered as both emitter and receiver of developmental regulatory signals having, on the one hand, informative and, on the other hand, structural roles. Thus, a central hypothesis in Developmental Biology proposes that the space-time organized operation of developmental cell behaviors depends on the cooperative establishment of spatially organized cell signaling networks mediated by diffusing informative molecules [3, 4, 5, 6]. Molecular diffusion results in asymmetric distribution of developmentally active informative signals. This asymmetry plays a fundamental role in establishing temporal and/or spatial organization of specific DCBs that result in the whole developmental process in the organized patterns of cells, tissues and organs, that living organisms eventually exhibit in their final, terminally differentiated, state.

### 1.1.2 The organizers and CNS patterning

Not all developing cell populations possess similarly relevant informative roles. There exist specific transient cell populations, the so-called organizers, that primarily play informative roles influencing or regulating the developmental behavior of the other cells. By means of installing asymmetric distributions of developmentally active signals the organizers serve as instruments of an informative reference system in the establishment of spatially organized processes of cell determination and differentiation [7, 8]. Amongst the most complex biological structures, the multilayered concentric neuronal organization of the central nervous system (CNS), i.e., brain cortex, cerebellum cortex etc., occupy a privileged position. The development of such a structural and functional complexity, the so-called corticogenesis, requires the organized operation of several DCBs. Amongst these DCBs: (a) the cell proliferation (CP), counteracted by apoptosis or programmed cell death, is involved in the generation of the appropriate number of neurons for each cortical area and each cortical layer; (b) the directed cell migration, a process mediated by specific interfacial interactions between cell surface and extracellular matrix components, controls the correct position of each specific neuronal type along the CNS spatial axes; (c) cell determination or commitment, a process mediated by irreversible genetic information reprogramming, allows different cell populations to select one out of a set of multiple developmental pathways; (d) cell differentiation, a process mediated by selective gene activation and selective protein synthesis, warrants the expression of different specific neuronal phenotypes; finally, (e) the development of the neural processes (neuritogenesis), dendrites and axons and the establishment of specific synaptic contacts (synaptogenesis) conclude this complex self-organizing and interactively regulated process of corticogenesis.

All these DCBs are properly organized in time and space. During the early stages, several organizers strategically positioned along the cephalic-caudal and dorsal-ventral axes of the developing CNS establish the primitive pattern of each region and sub-region, as well as the identity of the neuromeres. Later, cell proliferation, migration and differentiation controlled by reciprocal interactions allow the whole system to expand and differentiate into several structurally and functionally integrated systems of interconnected neuronal circuits. Abundant evidence indicate that all these processes are spatially and temporally organized [9, 10, 11, 12, 13, 14].

### 1.1.3 Roles of neuroepithelial cell proliferation and neuronal migration in corticogenesis

During the early development, the primitive CNS primordium, the so-called neural tube (NT), is almost exclusively composed of proliferative neuroepithelial (NE) cells. During this early proliferative phase the NE cells behave as a population of self-renewing stem cells that divide symmetrically, i.e. from each dividing NE cell originates two similar NE cells. These dividing NE cells are typically located along the inner surface of the NT forming the so-called generation zone (GZ). This type of early proliferation produces an amplification of the NE cells: they significantly increase in number, and the GZ increases in both area (planar expansion) and thickness (radial expansion). During a later proliferative phase NE cells proliferate asymmetrically, i.e. from each dividing NE cell originates a NE cell and a postmitotic neuron (PN). The number of NE cells and the GZ size stabilize while postmitotic neurons accumulate at an underlying premigratory zone (PMZ). Later, neurons migrate radially and the PMZ weakens while different layers of concentrically organized neurons begin to organize the future cortex. By the end of the proliferative phase the number of NE cells decreases, the GZ becomes thinner and finally disappears.

### 1.1.4 Neuroepithelial cell proliferation analyzed as a stochastic point process

The NE cells compose a synchronized and functionally integrated population of stem cells. Biomathematical analyses of numerical series representing the position of mitotic NE cells along the OT cephalic-caudal axis, analyzed within the framework provided by the theory of stochastic point processes, show that NE cells do not proliferate independently. It can be shown that the proliferative dynamics embeds (subsumes) two different components: (a) a non-stationary component that can be deterministically described and reveals a spatially organized process and (b) a stationary stochastic point process representing a uniformly distributed basal proliferative activity [15, 16]. More recently a different approach allows demonstrating that the stochastic component corresponds to an anti-correlated stationary process implying the result that there are short-range inhibitory interactions between neighboring proliferating NE cells [17, 18]. These mathematical results are consistent with experimental evidence indicating that cell proliferation is a DCB submitted to cell-cell controlling interactions and also to intracellular controlling processes.

### 1.1.5 Extra- and intracellular signals controlling proliferation dynamics

A general notion establishes that the coordinated behavior shown by proliferating cells during development [19] depends, on one hand, on a set of extracellular signals, the so-called growth factors [20, 21] and a set of intracellular informative signals composing a cell cycle control center whose main members are the regulatory proteins cyclins as well as cyclin-dependent kinases [22].

The temporally organized operation of these regulatory proteins installs the so-called cell division cycle (proliferative cycle). The cell division cycle is composed of a periodic ordered sequence of cyclic intracellular molecular events with typical durations. It involves two main phases: (a) the mitotic or M phase that includes the karyokinesis (nuclear division) and the cytokinesis (cytoplasm division) and (b) the interphase, or intermitotic phase. In turn, each of these phases involves temporally ordered sub-phases. The M phase is composed of the pro-, prometa-, meta-, ana- and telo-phase while the interphase is composed of the G1, S and G2 phases. Each type of proliferative cells population displays typical cell cycle length oscillating around a mean value. The total cell cycle length and also those of the different phases, vary,

depending on the cell type. Besides, particular cell types may change their cell cycle durations when long time series of subsequent periods are considered.

## 1.2 Biological settings of the present study

### 1.2.1 The experimental model system: NE cell proliferation in the developing avian optic tectum

The avian optic tectum (OT) is a mesencephalic alar plate derivative. The main OT input proceeds from the retinal ganglion cell axons. This information is integrated into others inputs, processed and transferred to other CNS areas. The OT possesses a typical cortical, multilayered concentric organization, consisting of alternating neuronal and fibrous layers. The developing avian OT is a well-characterized experimental model of CNS development. It is known that the OT determination and the positional information specifying the OT patterning are submitted to the organizing influence of the Isthmic Organizer (IsO). The IsO organizing activity is mediated by informative molecules which specify positional information along the OT cephalic-caudal axis [9, 10, 11, 12]. The positional information is firstly established as asymmetric gene expression and is then “translated”, by the differential operation of several DCBs into a gradient of cytoarchitectonic differentiation. There are well-documented data about its proliferative kinetics [23, 15, 16, 18], the postmitotic neuronal migration [15] and the expression of enzymes involved in neuronal migration [11], the histogenesis of its multilaminated architectural pattern [10] and also on the temporal and spatial developmental pattern of its specific neuronal types [13, 24]. There are also detailed structural, pharmacological and molecular studies on the development of its extrinsic innervation (serotonin innervation, retino-ectal connections) [9, 25, 26]. The developmental patterns of expression of several molecular components involved in synaptogenesis of the OT local circuits are also well characterized [14, 27, 28].

### 1.2.2 Biological methods and cell proliferation records

Empirical data on NE cell proliferation analyzed in this paper were recorded from complete histological serial sections obtained from OT of different developmental stages. OTs obtained from 2, 4 and 6 embryonic (E) days (*E2*, *E4*, *E6*) were used. During this interval the OT undergoes significant changes in size and shape. After dissection, OTs were processed for conventional histological methods [10]. Specimens were spatially oriented in order to obtain planar sections fulfilling criteria of adequacy between planar section orientation and developmental gradient axis position. These criteria were given in [6]. Cell proliferation records (CPR) along the OT cephalic-caudal axis were constructed on a video camera screen (Axioplan 2 imaging optical epifluorescence microscope coupled to an Axiocam HR color digital scanner and a computer equipped with the software Axiovision (all Carl Zeiss, Germany)) at a final magnification of 1000 X. A CPR is defined as a numerical sequence that indicates the density of mitotic cells in successive ordered,  $25\mu m$  length windows, along OT cephalic-caudal axis. The total number of cells in the GZ was also computed for each  $25\mu m$  window.

## 2 Complete mathematical model

The present modeling concerns the development of the OT between embryonic days *E2* and *E6*. It only describes the elongation of the cephalic-caudal axis and thus is only one space dimension. It also assumes that:

(a) the NE cells divide symmetrically (the number of asymmetric division is negligible),

- (b) the number of postmitotic neurons in the GZ is negligible and the increase in cell number in the GZ is mainly due to the symmetric divisions and
- (c) the cell cycle length does not change significantly neither along the interval  $E2$ - $E4$  nor between  $E4$  and  $E6$ . Consequently the numerical analysis and experimental validation treat separately these two periods.

We follow a classical description of the cell cycle that can be found in [29] and references therein. Our model is based on equations for the cell population densities structured by their progression along each phase  $0 \leq a \leq 1$  and their position  $x$  along the cephalic-caudal axis. The cells are divided in subclasses, as follows:

$n_1(t, a, x)$  = linear density of neuroepithelial cells in phases  $G_1, S, G_2$ ,

$n_2(t, a, x)$  = linear density of neuroepithelial cells in phase  $M$ ,

$n_3(t, a, x)$  = postmitotic neurons generated by asymmetric mitosis. These cells stop proliferating; they are destined to migrate and later differentiate into specialized neurons.

We propose a unique model for time  $t$  between days  $E2$  and  $E10$ . Cells in the phases are also subject to random motion along the cephalic-caudal axis and this leads us to represent the dynamics of cell division and motion by the equations

$$\begin{cases} \frac{\partial n_1}{\partial t} + \frac{\partial[v_1(t)n_1]}{\partial a} = \frac{\partial^2[D(x)n_1]}{\partial x^2}, \\ \frac{\partial n_2}{\partial t} + \frac{\partial[v_2(t)n_2]}{\partial a} = \frac{\partial^2[D(x)n_2]}{\partial x^2}, \\ \frac{\partial n_3}{\partial t} = q(t, x)v_2(t)n_2(t, a = 1, x). \end{cases} \quad (1)$$

This system is completed with boundary and initial conditions

$$\begin{cases} v_2(t)n_2(t, a = 0, x) = v_1(t)n_1(t, a = 1, x), \\ v_1(t)n_1(t, a = 0, x) = [2 - q(t, x)]v_2(t)n_2(t, a = 1, x), \\ \frac{\partial n_1}{\partial x}(t, a, x = x_+(t)) = 0, \quad \frac{\partial n_2}{\partial x}(t, a, x = x_+(t)) = 0, \quad \frac{\partial n_3}{\partial x}(t, a, x = x_+(t)) = 0, \\ n_1(0, a, x) = n_1^0(a, x), \quad n_2(0, a, x) = n_2^0(a, x), \quad n_3(0, a, x) = 0. \end{cases} \quad (2)$$

We have used the following notations

- $t$  the chronological time variable,
- $x$ : abscissa of transverse sections along the cephalic-caudal OT axis,  $0 \leq x \leq x_+(t)$ ,  $x = 0$  cephalic tip,  $x = x_+(t)$ , caudal tip at time  $t$ ,
- $0 \leq a \leq 1$ : degree of progression along the mitotic ( $M$ ) phase or the intermitotic ( $G_1, S, G_2$ ) phase,
- $v_i(t)$ , ( $i = 1, 2$ ), homogeneous to an inverse of time, is the frequency of the corresponding phase duration,
- $D(x)$ : “effective cellular diffusion” at section  $x$ ,
- $0 \leq q(t, x) \leq 1$ : postmitotic neuronal population growth rate, or rate of differentiation (a more complete model should consider it as a nonlinear function of the cell densities).

The model should also be completed with a law for the OT elongation along the development, which means a description of the  $x_+(t)$ . It should naturally take the form

$$\frac{d}{dt}x_+(t) = F(n_1(t), n_2(t), n_3(t)), \quad (3)$$

with a function  $F(\cdot)$  that describes how the OT development is ruled by the different cell types. In this case the Neumann boundary conditions at  $x^+(t)$  should be replaced by the no-flux conditions

$$\dot{x}_+(t)n_i(t, a, x_+(t)) + D(x)\frac{\partial}{\partial x}n_i(t, a, x = x_+(t)) = 0.$$

To access such nonlinear coefficients as  $q(\cdot)$  and  $F(\cdot)$  is too demanding in a first stage, and thus simplifications are needed. Therefore we reduce the model complexity with two assumptions. Firstly, for days between  $E2$  and  $E6$  we can neglect asymmetric cell proliferation which leads to  $q = 0$  and  $n_3 = 0$ . Secondly, we assume that the OT has constant length between days  $E2$ - $E4$  and  $E4$ - $E6$ ; this leads to fix  $x_+$  independent of time with a jump at the end of day  $E4$ .

### 3 Reduced model for the symmetric proliferative phase without postmitotic migrating neurons ( $q = 0$ )

We reduce the model complexity in the present work with two assumptions. Firstly, for days between  $E2$  and  $E6$  we can neglect asymmetric cell proliferation which leads to  $q = 0$  and  $n_3 = 0$ . Secondly, we assume that the OT has constant length between days  $E2$ - $E4$  and  $E4$ - $E6$ ; this leads to fix  $x_+$  independent of time with a jump at the end of day  $E4$ . According to these assumptions, we consider only a simplified version of the model that we present now. We also present its discretization.

#### 3.1 The simplified model

We define the region

$$\Omega := \{0 \leq t \leq T, \quad 0 \leq a \leq 1, \quad 0 \leq x \leq x_+\},$$

with  $x_+$  constant and we also suppose that  $v_1, v_2, D$  constant. This leads to the model defined by the system of partial differential equations

$$\begin{cases} \frac{\partial n_1}{\partial t} + v_1 \frac{\partial n_1}{\partial a} = D \frac{\partial^2 n_1}{\partial x^2}, \\ \frac{\partial n_2}{\partial t} + v_2 \frac{\partial n_2}{\partial a} = D \frac{\partial^2 n_2}{\partial x^2}, \end{cases} \quad (4)$$

completed with the boundary and initial conditions

$$\begin{cases} v_2 n_2(t, a = 0, x) = v_1 n_1(t, a = 1, x), \\ v_1 n_1(t, a = 0, x) = 2v_2 n_2(t, a = 1, x), \\ \frac{\partial n_1}{\partial x}(t, a, x = x_+(t)) = 0, \quad \frac{\partial n_2}{\partial x}(t, a, x = x_+(t)) = 0, \\ n_1(0, a, x) = n_1^0(a, x), \quad n_2(0, a, x) = n_2^0(a, x). \end{cases} \quad (5)$$

For later purposes we impose the compatibility condition between the boundary and initial conditions:

$$v_1 n_1^0(1, x) = v_2 n_2^0(0, x) \quad (6)$$

It implies that the solution is smooth enough for our analysis below, see [5].

#### 3.2 Discretization by the method of finite differences

For every  $K \in \mathbb{N}$ ,  $I \in \mathbb{N}$ ,  $J \in \mathbb{N}$  we define a mesh in  $\Omega$  by the points

$$(k\Delta x, i\Delta a, j\Delta t), \quad k = 0, 1, \dots, K, \quad i = 0, 1, \dots, I, \quad j = 0, 1, \dots, J$$

with

$$\Delta t = \frac{T}{K}, \quad \Delta a = \frac{1}{I}, \quad \Delta x = \frac{x_+}{J}.$$

For explicit schemes, these parameters have to be chosen with the stability (Courant-Friedrichs-Lewy) condition

$$v \frac{\Delta t}{\Delta a} + 2D \frac{\Delta t}{(\Delta x)^2} \leq 1. \quad (7)$$

According to well established methods [30, 31], we approximate the derivatives in the differential equations by upwind (for first order derivatives) or centered (for the second order derivatives) differences on the mesh points. For a generic point of the mesh  $(k\Delta x, i\Delta a, j\Delta t)$  we set as usual

$$\begin{aligned} \frac{\partial n_l}{\partial t} &\cong \frac{n_l(k+1, i, j) - n_l(k, i, j)}{\Delta t}, & \frac{\partial n_l}{\partial a} &\cong \frac{n_l(k, i, j) - n_l(k, i-1, j)}{\Delta a} \\ \frac{\partial^2 n_l}{\partial x^2} &\cong \frac{n_l(k, i, j+1) - 2n_l(k, i, j) + n_l(k, i, j-1)}{(\Delta x)^2}, & l &= 1, 2. \end{aligned}$$

In the following we will denote by:

- $n(k, i, j)$  value of the exact solution at the point  $(k\Delta x, i\Delta a, j\Delta t)$  of the mesh,
  - $u(k, i, j)$  value of the discretized solution at the point  $(k)\Delta x, i\Delta a, j\Delta t$ ,
- and we consider the system of equations obtained upon application of these approximations, with the corresponding boundary and initial conditions.

The discretized model reads, for  $i = 1, \dots, I, j = 1, \dots, J-1, k = 0, 1, \dots, K$ ,

$$\left\{ \begin{aligned} \frac{u_1(k+1, i, j) - u_1(k, i, j)}{\Delta t} + v_1 \frac{u_1(k, i, j) - u_1(k, i-1, j)}{\Delta a} &= D \frac{u_1(k, i, j+1) - 2u_1(k, i, j) + u_1(k, i, j-1)}{(\Delta x)^2}, \\ \frac{u_2(k+1, i, j) - u_2(k, i, j)}{\Delta t} + v_2 \frac{u_2(k, i, j) - u_2(k, i-1, j)}{\Delta a} &= D \frac{u_2(k, i, j+1) - 2u_2(k, i, j) + u_2(k, i, j-1)}{(\Delta x)^2}, \\ v_2 u_2(k, 0, j) &= v_1 u_1(k, I, j), \\ v_1 u_1(k, 0, j) &= 2v_2 u_2(k, I, j), \\ u_1(0, i, j) &= n_1^0(i, j), & u_2(0, i, j) &= n_2^0(i, j). \end{aligned} \right. \quad (8)$$

It is again convenient to also assume a compatibility condition between the boundary and initial conditions in the discretized model

$$v_1 n_1^0(I, j) = v_2 n_2^0(0, j) \quad j = 0, 1, \dots, J. \quad (9)$$

We use  $u$  to indicate both  $u_1$  and  $u_2$ , and  $v$  for  $v_1$  and  $v_2$ . Therefore, from equation (8), we deduce

$$\begin{aligned} \frac{u(k+1, i, j) - u(k, i, j)}{\Delta t} + v \frac{u(k, i, j) - u(k, i-1, j)}{\Delta a} &= \\ &= D \frac{u(k, i, j+1) - 2u(k, i, j) + u(k, i, j-1)}{(\Delta x)^2}. \end{aligned}$$

In other words, we obtain  $u(k+1, i, j)$  thanks to the iterations on the time label

$$\begin{aligned} u(k+1, i, j) &= u(k, i, j) - v \frac{\Delta t}{\Delta a} (u(k, i, j) - u(k, i-1, j)) + \\ &+ D \frac{\Delta t}{(\Delta x)^2} (u(k, i, j+1) - 2u(k, i, j) + u(k, i, j-1)). \end{aligned} \quad (10)$$

We use the discrete Neumann boundary conditions

$$u(k, i, 0) = u(k, i, 1), \quad u(k, i, J+1) = u(k, i, J), \quad (11)$$

for  $k = 0, 1, \dots, K, i = 0, 1, \dots, I$ .

Applying the induction formula (10) to calculate the values of  $u_1$  and  $u_2$ , and bearing (11) and the initial conditions in mind, we obtain the values of the discretized solution

$$u_1(k, i, j), \quad u_2(k, i, j), \quad k = 0, 1, \dots, K, \quad i = 0, 1, \dots, I, \quad j = 0, 1, \dots, J. \quad (12)$$

This describes the numerical method we use in the sequel together with experimental data.

### 3.3 Convergence of the approximate solutions to the exact solution on the points of the mesh

The convergence analysis of this numerical scheme is standard and can be found in [30, 31]. We give here a fast account for the sake of completeness.

We use the notation  $n(k, i, j)$  to indicate both  $n_1(k, i, j)$  and  $n_2(k, i, j)$ , and also  $u(k, i, j)$  for  $u_1(k, i, j)$  and  $u_2(k, i, j)$ .

Applying Taylor's formula to the exact solution at the points of the mesh we obtain:

$$\begin{aligned} \frac{n(k+1, i, j) - n(k, i, j)}{\Delta t} + v \frac{n(k, i, j) - n(k, i-1, j)}{\Delta a} = \\ = D \frac{n(k, i, j+1) - 2n(k, i, j) + n(k, i, j-1)}{(\Delta x)^2} + \Psi, \end{aligned} \quad (13)$$

where for some constants  $A, B, C$ , we have

$$|\Psi| \leq A\Delta t + B\Delta a + C(\Delta x)^2. \quad (14)$$

From (13) we obtain

$$\begin{aligned} n(k+1, i, j) = & n(k, i, j) - v \frac{\Delta t}{\Delta a} (n(k, i, j) - n(k, i-1, j)) + \\ & + D \frac{\Delta t}{(\Delta x)^2} (n(k, i, j+1) - 2n(k, i, j) + n(k, i, j-1)) + \Psi \Delta t. \end{aligned} \quad (15)$$

Next, we denote the error introduced by approximating  $n(k, i, j)$  by  $u(k, i, j)$  as

$$w(k, i, j) = n(k, i, j) - u(k, i, j) \quad (16)$$

From (10) and (15), we obtain

$$\begin{aligned} w(k+1, i, j) = & \left(1 - v \frac{\Delta t}{\Delta a} - 2D \frac{\Delta t}{(\Delta x)^2}\right) w(k, i, j) + v \frac{\Delta t}{\Delta a} w(k, i-1, j) \\ & + D \frac{\Delta t}{(\Delta x)^2} (w(k, i, j+1) + w(k, i, j-1)) + \Psi \Delta t. \end{aligned} \quad (17)$$

From the stability condition (7) and using (17), we obtain

$$\begin{aligned} |w(k+1, i, j)| \leq & \left(1 - v \frac{\Delta t}{\Delta a} - 2D \frac{\Delta t}{(\Delta x)^2}\right) |w(k, i, j)| + v \frac{\Delta t}{\Delta a} |w(k, i-1, j)| + \\ & + D \frac{\Delta t}{(\Delta x)^2} (|w(k, i, j+1)| + |w(k, i, j-1)|) + |\Psi| \Delta t \end{aligned} \quad (18)$$

For every  $k$  we define

$$M_k = \max_{i,j} |w(i, j, k)|.$$

From (18), we obtain successively

$$M_{k+1} \leq \left(1 - v \frac{\Delta t}{\Delta a} - 2D \frac{\Delta t}{(\Delta x)^2}\right) M_k + v \frac{\Delta t}{\Delta a} M_k + 2D \frac{\Delta t}{(\Delta x)^2} M_k + |\Psi| \Delta t,$$

$$M_{k+1} \leq M_k + |\Psi| \Delta t.$$

Summing over  $k = 0, 1, \dots, m-1$ ,  $m \leq K$

$$\begin{aligned} \sum_{k=0}^{m-1} M_{k+1} & \leq \sum_{k=0}^{m-1} M_k + m\Delta t |\Psi|, \\ M_m & \leq M_0 + m\Delta t |\Psi|. \end{aligned} \quad (19)$$

Let us observe that  $u(0, i\Delta a, j\Delta x) = n(0, i\Delta a, j\Delta x)$ , and therefore we have

$$w(0, i, j) = w(0, i\Delta a, j\Delta x) = 0, \quad (i = 0, 1, \dots, I, \quad j = 0, 1, \dots, J).$$



Bearing in mind that  $M_0 = 0$  and  $\Delta t = \frac{T}{K}$ , from (19) we obtain

$$M_m \leq m \frac{T}{K} |\Psi|.$$

Since  $\frac{m}{K} \leq 1$ , we also conclude that  $M_m \leq |\Psi|T$ .

On the other hand, from (14), we finally obtain

$$M_m \leq (A\Delta t + B\Delta a + C(\Delta x)^2) T, \quad m = 0, 1, \dots, K.$$

This proves that

$$\max_{k,i,j} |n(k,i,j) - u(k,i,j)| \rightarrow 0 \quad \text{as} \quad \Delta t \rightarrow 0, \Delta a \rightarrow 0, \Delta x \rightarrow 0.$$

The previous reasoning is valid for both components of the solution of the system (4), therefore

$$\max_{k,i,j} |n_1(k,i,j) - u_1(k,i,j)| \rightarrow 0 \quad \text{and} \quad \max_{k,i,j} |n_2(k,i,j) - u_2(k,i,j)| \rightarrow 0,$$

as  $\Delta t \rightarrow 0, \Delta a \rightarrow 0, \Delta x \rightarrow 0$ .

## 4 Dynamics of the simplified model from $E2$ to $E4$

We now study the dynamics of the model between  $E2$  and  $E4$ . It is assumed that only symmetric mitosis takes place during that period. Then, we use the model (discretized by finite differences) given by equations (8).

### 4.1 Experimental data recorded from the optic tectum at $E2$

Records of number of cells counts performed on successive segments along the optic tectum cephalic-caudal axis were used as initial data for the model. Such information at day  $E2$  corresponds to 16 segments,  $25\mu m$ . We have named these segments "sections". Every segment is identified by a value of the abscissa  $x$ .

The initial model length is  $L_{E2} = 16 \times 25\mu m = 400\mu m$ , and the total initial cells number is  $N_{E2} = 800$ .

The distribution of cells number per section is reported in TABLE 1 (Annexe)

### 4.2 Determination of parameters

**Cell cycle duration between  $E2$  and  $E4$ .** According to experimental data, in the interval  $E2 - E4$ , the number of cells increases from approximately around 800 to around 9200 cells. Assuming that during this period the whole neuro-epithelial cells population exhibits a homogeneous proliferative behavior with a cell cycle duration around a mean value, then the averaged duration can be estimated as 13.6 hours.

- Cell cycle duration:  $G_1 + S + G_2 + M = 13h36min = 13.6h$ .
- Intermitotic cycle duration  $G_1 + S + G_2$  is 96 % the cell cycle duration; hence,
- Duration of  $G_1 + S + G_2 = 13.056h \cong 13h03min$ .

The mitotic cycle duration  $M$  represents 4% the cell cycle duration :

- Mitotic duration  $M = 0.544h = 33min$ .

**Number of cycles  $G_1 + S + G_2 + M$  between  $E2$  and  $E4$ .**

$$n_{E2-4} = \frac{48hs}{13.6hs} \cong 3.53$$

**Total number of cells at  $E4$ .** Using the formula

$$N_{E4} = N_{E2} \times 2^{n_{E2-4}}$$

we obtain

$$N_{E4} = 800 \times 2^{3.53} \cong 9241$$

**Cellular density.** It was observed experimentally that the average cellular density  $\delta_{E2-4}$  between  $E2$  and  $E4$  remained approximately constant,  $3.85 \frac{cel}{\mu m}$ . Then, it will be assumed that between  $E2$  and  $E4$  the model average cell density corresponds to this value.

$$\delta_{E2-4} = 3.85 \frac{cel}{\mu m}$$

**Model length at  $E4$ .**

$$L_{E4} = \frac{N_{E4}}{\delta_{E2-4}} = \frac{9241 cel}{3.85 \frac{cel}{\mu m}} \cong 2400 \mu m$$

**Number of  $25\mu m$  thickness sections comprising the model at  $E4$ .**

$$D_{E2-4} \cong \frac{2400 \mu m}{400 \mu m} = 6$$

**Calculation of  $v_1$  and  $v_2$  between  $E2$  and  $E4$ .** The following formulae define  $v_1$  and  $v_2$ ,

- $\frac{1}{v_1} = n^{er}$  of interphases  $G_1 + S + G_2$  per day =  $\frac{24hs}{interphase\ duration\ in\ hs}$
- $\frac{1}{v_2} = n^{er}$  of mitosis  $M$  by day =  $\frac{24hs}{mitotic\ duration\ in\ hs}$

In our case we have

$$\frac{1}{v_1} \cong \frac{24}{13} \cong 1.85 \quad \frac{1}{v_2} \cong \frac{24}{0.544} \cong 44.12$$

and we obtain

$$v_1 \cong 0.54, \quad v_2 \cong 0.023.$$

**Time interval.** We take  $t = 0$  at  $E2$  and  $t = T = 48hs$  at  $E4$ . We decompose the interval in sub-intervals of 10 minutes and we take  $\Delta t = 10min = \frac{1}{6}hour$ . The index will be  $k = 0, 1, \dots, 48 \times 6 = 288$ ,  $K = 288$ .

**Interval on the longitudinal axis.** We decompose the interval  $[0, 400]$  in 16 sub-intervals of length  $\Delta x = 25\mu m$ .

The index will be  $j = 0, 1, \dots, 16$   $J = 16$ .

**Interval of variation of  $a$ .** The interval of variation of  $a$  is  $[0, 1]$ . We will choose  $\Delta a$  to satisfy the stability condition

$$v_1 \frac{\Delta t}{\Delta a} + 2D \frac{\Delta t}{(\Delta x)^2} \leq 1, \quad v_2 \frac{\Delta t}{\Delta a} + 2D \frac{\Delta t}{(\Delta x)^2} \leq 1.$$

Since

$$v_1 = 0.54 \frac{1}{h}, \quad \Delta t = \frac{1}{6}h, \quad D = 6 \frac{(\mu m)^2}{h}, \quad \Delta x = 25\mu m, \quad v_2 = 0.023 \frac{1}{h} \quad (20)$$

the condition of stability will be the following one:

$$0.54 \frac{1/6}{\Delta a} + 12 \frac{1/6}{625} \leq 1, \quad 0.023 \frac{1/6}{\Delta a} + 12 \frac{1/6}{625} \leq 1.$$

From the stability condition (7), it is sufficient to choose  $\Delta a$  such that

$$0.54 \frac{1/6}{\Delta a} + 12 \frac{1/6}{625} \leq 1, \quad or \quad \Delta a > 0.09.$$

We will adopt that index will be  $i = 0, 1, \dots, 10$  that is

$$I = 10, \quad \Delta a = 0.1. \quad (21)$$

### 4.3 Induction formula for $u_1$

$$\begin{aligned} u_1(k+1, i, j) &= u_1(k, i, j) - v_1 \frac{\Delta t}{\Delta a} (u_1(k, i, j) - u_1(k, i-1, j)) + \\ &\quad + D \frac{\Delta t}{(\Delta x)^2} (u_1(k, i, j+1) - 2u_1(k, i, j) + u_1(k, i, j-1)). \end{aligned} \quad (22)$$

Summing over  $i$  from  $i=0$  to  $i=I$  in (19), we obtain

$$\begin{aligned} \sum_{i=0}^I u_1(k+1, i, j) &= \sum_{i=0}^I u_1(k, i, j) - v_1 \frac{\Delta t}{\Delta a} \left\{ \sum_{i=0}^I u_1(k, i, j) - \sum_{i=0}^I u_1(k, i-1, j) \right\} + \\ &\quad + D \frac{\Delta t}{(\Delta x)^2} \left\{ \sum_{i=0}^I u_1(k, i, j+1) - 2 \sum_{i=0}^I u_1(k, i, j) + \sum_{i=0}^I u_1(k, i, j-1) \right\}. \end{aligned} \quad (23)$$

Observe that

$$\sum_{i=0}^I u_1(k, i, j) - \sum_{i=0}^I u_1(k, i-1, j) = u_1(k, I, j) - u_1(k, -1, j),$$

therefore, from (23) we obtain

$$\begin{aligned} \sum_{i=0}^I u_1(k+1, i, j) &= \sum_{i=0}^I u_1(k, i, j) - v_1 \frac{\Delta t}{\Delta a} \{u_1(k, I, j) - u_1(k, -1, j)\} + \\ &\quad + D \frac{\Delta t}{(\Delta x)^2} \left\{ \sum_{i=0}^I u_1(k, i, j+1) - 2 \sum_{i=0}^I u_1(k, i, j) + \sum_{i=0}^I u_1(k, i, j-1) \right\}. \end{aligned} \quad (24)$$

Let us also observe that the sum

$$U_1(l, r) = \sum_{i=0}^I u_1(l, i, r)$$

is the total number of interphasic cells ( $G1 + S + G2$ ) at instant  $l$  and at section  $r$ . Replacing in (24) we obtain

$$\begin{aligned} U_1(k+1, j) &= U_1(k, j) - v_1 \frac{\Delta t}{\Delta a} \{u_1(k, I, j) - u_1(k, -1, j)\} + \\ &\quad + D \frac{\Delta t}{(\Delta x)^2} \{U_1(k, j+1) - 2U_1(k, j) + U_1(k, j-1)\}. \end{aligned} \quad (25)$$

$u_1(k, I, j)$  is the number of cells that complete the interphase  $G1 + S + G2$  at instant  $k$ , section  $j$ .

$u_1(k, 0, j)$  is the number of cells that begin the interphase  $G1 + S + G2$  at instant  $k$ , section  $j$ .

The number  $u_1(k, 0, j) - u_1(k, I, j)$  depends on the total number of interphasic cells  $G1 + S + G2$ , at instant  $k$  and section  $j$ ; therefore, it is a function of  $U_1(k, j)$ .

It is difficult establishing this dependence law. It will be assumed that there exist a constant  $\alpha(k, j)$ , that depends on instant  $k$  and section  $j$ , such that

$$u_1(k, 0, j) - u_1(k, I, j) = \alpha(k, j) U_1(k, j). \quad (26)$$

Now we replace in (25), bearing in mind (26) and the values of  $v_1, \Delta t, \Delta x, \Delta a, D$  in (20), to obtain

$$U_1(k+1, j) = (1 + 0.9\alpha(j))U_1(k, j) + 0.0032 \left( \frac{U_1(k, j+1) + U_1(k, j-1)}{2} - U_1(k, j) \right). \quad (27)$$

### 4.4 Approximation of the induction formulæ

Since, for each  $k$ , the function  $U_1(k, j)$  is continuous in  $j$ , we conclude that

$$0.0032 \left( \frac{U_1(k, j+1) + U_1(k, j-1)}{2} - U_1(k, j) \right)$$

will be very small. Therefore, it will be ignored in a first approximation. Then, the formula (27) takes the following approximate form

$$U_1(k+1, j) = (1 + 0.9\alpha(j)) U_1(k, j). \quad (28)$$

Multiplying expressions (28) for  $k = 0, 1, \dots, 287$  we obtain

$$\begin{aligned} \prod_{k=0}^{287} U_1(k+1, j) &= (1 + 0.9\alpha(j))^{288} \prod_{k=0}^{287} U_1(k, j), \quad \text{or} \\ U_1(48hs, j) &= (1 + 0.9\alpha(j))^{288} U_1(0, j). \end{aligned} \quad (29)$$

Now,  $U_1(48hs, j)$  represents the number of cells generated between  $E2$  and  $E4$  by the cells that at  $E2$  were located at segment  $j$ . Given that during this period the model length increases 6 fold, it is assumed that, at  $E4$  these newly generated cells will expand over six  $25\mu m$  thickness segments.

Thus, cells located at segment at  $E2$  give origin to cells that at  $E4$  occupy  $s$  - sections from  $s = 1 + 6(j-1)$  to  $s = 6j$ . The values of  $j$  will be  $j = 1, 2, \dots, 16$ , corresponding to 16 sections of the model at  $E2$ .

$U_1(48hs, s)$  indicates the number of cells located at segment of abscissa  $s$  at  $E4$ .

$\bar{U}_1(48hs, s)_{s=1+6(j-1)}^{s=6j}$  will indicate the average number of cells in the segments of abscissa  $s$  at  $E4$ , for  $1 + 6(j-1) \leq s < 6j$ .

We have

$$\bar{U}_1(48hs, s)_{s=1+6(j-1)}^{s=6j} = \frac{U_1(48hs, j)}{6}.$$

For the sake of brevity, we will sometimes write  $\bar{U}_1(48hs, s)$  instead of  $\bar{U}_1(48hs, s)_{s=1+6(j-1)}^{s=6j}$  for the corresponding value of  $j$  which appears in the above mentioned formula. We will have

$$\begin{aligned} \bar{U}_1(48hs, s) &= \frac{1}{6}(1 + 0.9\alpha(j))^{288} U_1(0, j), \quad 1 + 6(j-1) \leq s < 6j, \quad \text{or} \\ \bar{U}_1(48hs, s) &= \beta(j) U_1(0, j), \quad j = 1, 2, \dots, 16, \end{aligned} \quad (30)$$

where

$$\beta(j) = \frac{1}{6} (1 + 0.9\alpha(j))^{288}.$$

We define the cell proliferation at segment  $j$  as the quotient

$$cel.prol.(j) = \frac{U_1(48hs, j)}{U_1(0hs, j)}.$$

Since  $U_1(48hs, j) = 6 \bar{U}_1(48hs, s)$ , we obtain

$$cel.prol.(j) = 6\beta(j).$$

$\beta(j)$  is a coefficient proportional to the cell proliferation at segment  $j$ .

The relationship between  $\alpha(j)$  and  $\beta(j)$  for  $j = 0, 1, \dots, 16$  is

$$\alpha(j) = \frac{1}{0.9} \left( [6\beta(j)]^{\frac{1}{288}} - 1 \right) \quad \beta(j) = \frac{1}{6} (1 + 0.9\alpha(j))^{288}$$

It will be assumed that proliferation is constant along the model, with a value equal to an average proliferation. This is equivalent to assume that  $\alpha(j) = \alpha$  and  $\beta(j) = \beta$  are independent of  $j$ .

Finally, we will have

$$\bar{U}_1(48hs, s) = \beta(j) U_1(0, j) \quad j = 1, 2, \dots, 16. \quad (31)$$

We will calculate the constants  $\alpha$  and  $\beta$ . We will consider a hypothetical segment  $j$  in which the number of cells will be obtained by calculating the average of the total number of cells of 16 sections in which the model was divided at  $E2$ . This number is

$$\frac{800}{16} = 50.$$

Likewise, we will consider that cells of section  $j$  originate a number of cells that at  $E4$  occupy 6 sections of the model. The number of cells at each one of the above mentioned sections is obtained by calculating the average of the totality of cells of the 96 sections comprising the model. This average value is

$$\frac{9241}{96} \cong 96.26$$

and, therefore, 578 cells correspond to 6 sections. We will apply the formula (29) taking

$$U_1(0, j) = 50 \quad U_1(48hs, j) = 578.$$

We obtain  
and therefore

$$578 = (1 + 0.9\alpha)^{288} 50$$

$$\alpha = \frac{10}{9} \left( \left[ \frac{578}{50} \right]^{\frac{1}{288}} - 1 \right).$$

We conclude that

$$\alpha \cong 0.0095 \quad \text{and} \quad \beta \cong 1.93 \quad (32)$$

Replacing in (31) we obtain

$$\bar{U}_1(48hs, s) = 1.93U_1(0, j) \quad 1 + 6(j-1) \leq s \leq 6j, \quad j = 1, 2, \dots, 16. \quad (33)$$

#### 4.5 Number of cells given by formula (30) and comparison with experimental data

The distribution of cells number per section is reported in TABLE 2 (Annexe)

#### 4.6 Approximation for linear interpolation of the number of cells in the intermediate sections and comparison with experimental data corresponding to $E4$

We will put

$$\bar{U}_1(s) = \bar{U}_1(48hs, s), \quad 1 + 6(j-1) \leq s \leq 6j, \quad j = 1, 2, \dots, 16.$$

We will use the formula of linear interpolation

$$\bar{U}_1(s) = \bar{U}_1(1 + 6(j-1)) + \frac{\bar{U}_1(1 + 6j) - \bar{U}_1(1 + 6(j-1))}{6}(s - 1) \quad (34)$$

for  $1 + 6(j-1) \leq s \leq 6j, j = 1, 2, \dots, 16$ .

The value  $\bar{U}_1(97)$  used in the calculation for the slope in interval  $91 \leq s \leq 96$  is obtained by means of the following trick. Let us observe that in the initial tip of the model at  $E2$  the variation in the cell number between segments  $j = 1$  and  $j = 2$  is, in increasing sense

$$U_1(0, j = 2) - U_1(0, j = 1) = 7$$

This induces us to extrapolate TABLE 2 by putting  $U_1(0, j = 17) = 42$ , after which the variation at the model final tip will be in diminishing sense, of the same magnitude,

$$U_1(0, j = 17) - U_1(0, j = 16) = -7$$

Applying formula (33) for  $j = 17$  we obtain

$$\bar{U}_1(s) = \bar{U}_1(48hs, s) \cong 1.93 \times 42 \cong 81 \quad \text{for } 97 \leq s \leq 102.$$

We take the value  $\bar{U}_1(97) = 81$ .

Total number of cells per section  $s$  at  $E4$  is reported in TABLE 3 (Annexe) and Fig. 1.

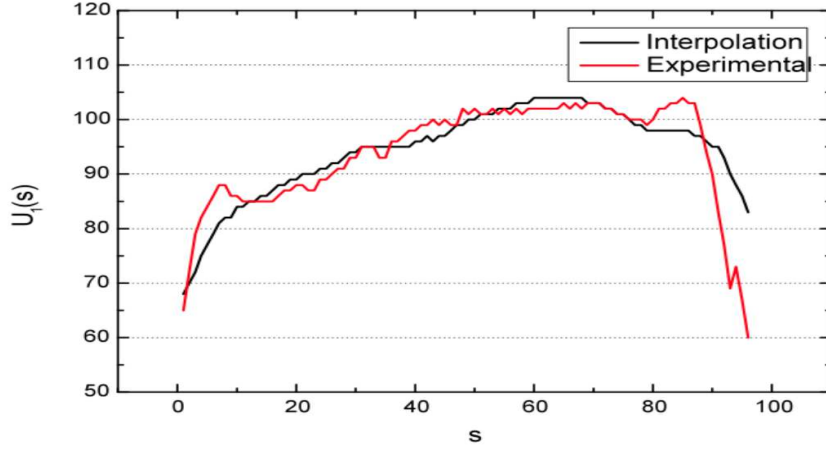


Figure 1: Total number of cells per section  $s$  in E4.

## 5 Dynamics of the model between $E4$ and $E6$

### 5.1 Initial data

The number of cells calculated by the model for an  $E4$  optic tectum was taken as initial data to calculate the evolution between  $E4$  and  $E6$  and to estimate the number of cell each section will originate over that period.

At  $E4$  the model is composed of 96 segments ,  $25\mu m$  thickness.  
The model initial length at  $E4$  is  $L_{E4} = 96 \times 25\mu m = 2400\mu m$   
The total initial number of cells of the model at  $E4$  is  $N_{E4} = 9241$ .

Table 3 illustrates spatially ordered distribution of number of cells/segment mathematically (left) and empirically (right) estimated.

### 5.2 Determination of parameters

**Cell cycle duration between  $E4$  and  $E6$ .** Based on assumptions introduced in a previous section (4.2.) the average duration of the complete cell cycle  $G_1 + S + G_2 + M$  can be estimated as around as around 38 hs 15 min.

Cell cycle duration  $G_1 + S + G_2 + M = 38.25h = 38h15min$ .

Assuming that the interphase  $G_1 + S + G_2$  is 96 % the complete cycle,

Interphase duration  $G_1 + S + G_2 = 36.72h \cong 36h43min$ .

The mitotic length  $M$  is 4 % the complete cycle length,  
Mitotic duration  $M = 4\%$  of  $38.25h = 1.53h \cong 1h32min$  .

**Number of cycles  $G_1 + S + G_2 + M$  between  $E4$  and  $E6$ .**

$$n_{E4-6} = \frac{48hs}{38.25hs} \cong 1.255$$

**Total number of cells of the model at  $E6$ .** Using the formula

$$N_{E6} = N_{E4} \times 2^{n_{E4-6}}$$

we obtain

$$N_{E6} = 9241 \times 2^{1.255} \cong 22055$$

**Cellular density.** It was experimentally observed that the average cell density  $\delta_{E4-6}$  of the Tectum between  $E4$  and  $E6$  remains approximately constant, with a value of  $3.85 \text{ cel}/\mu m$ .

We will suppose therefore that  $\delta_{E4-6} = 3.85 \text{ cel}/\mu\text{m}$ .

**Model length at E6.** We have  $L_{E6} = \frac{N_{E6}}{\delta_{E4-6}} = \frac{22055 \text{ cel}}{3.85 \text{ cel}/\mu\text{m}} \cong 5729 \mu\text{m}$

**Number of  $25\mu\text{m}$  thickness segments comprising the model at E6.**

$$n^{er} \text{ of segments} = \frac{L_{E6}}{25\mu\text{m}} = \frac{5729\mu\text{m}}{25\mu\text{m}} \cong 230$$

**Coefficient of cell diffusion between E4 and E6.**

$$D_{E4-6} = \frac{5729\mu\text{m}}{2400\mu\text{m}} = 2.387... \cong 2.4\mu\text{m}$$

**Calculation of  $v_1$  and  $v_2$  between E4 and E6.** The formulae that define  $v_1$  and  $v_2$  are the following ones

- $\frac{1}{v_1} = n^{er}$  of interphases  $G_1 + S + G_2$  per day =  $\frac{24\text{hs}}{\text{interphase duration in hs}}$
- $\frac{1}{v_2} = n^{er}$  of mitosis  $M$  per day =  $\frac{24\text{hs}}{\text{mitosis duration in hs}}$

We have  $\frac{1}{v_1} = \frac{24}{36.72}$  and  $\frac{1}{v_2} = \frac{24}{1.53}$  therefore

$$v_1 = \frac{36.72}{24} = 1.53, \quad v_2 = \frac{1.53}{24} \cong 0.064.$$

**Time interval.** We take  $t = 0$  at E4 and  $t = T = 48\text{hs}$  at E6. We decompose the interval in sub-intervals of 10 minutes each and we take

$$\Delta t = 10 \text{ min} = \frac{1}{6} \text{ hour}.$$

The index will be  $k = 0, 1, \dots, 48 \times 6 = 288$ ,  $K = 288$ .

**Interval on the longitudinal axis.** We decompose the interval  $[0, 2400]$  in 96 sub-intervals of length  $\Delta x = 25\mu\text{m}$ . The index will be  $j = 0, 1, \dots, 96$ ,  $J = 96$ .

**Interval of variation of  $a$ .** The interval of variation for  $a$  is still  $[0, 1]$ . We choose  $\Delta a$  so that the stability condition (7) is satisfied

$$\nu_1 \frac{\Delta t}{\Delta a} + 2D \frac{\Delta t}{(\Delta x)^2} \leq 1 \quad \nu_1 \frac{\Delta t}{\Delta a} + 2D \frac{\Delta t}{(\Delta x)^2} \leq 1.$$

Since

$$v_1 = 1.53 \frac{1}{h}, \Delta t = \frac{1}{6} h, D = 2.4 \frac{(\mu\text{m})^2}{h}, \Delta x = 25\mu\text{m}, v_2 = 0.064 \frac{1}{h}. \quad (35)$$

the stability condition becomes

$$1.53 \frac{1/6}{\Delta a} + 4.8 \frac{1/6}{625} \leq 1, \quad 0.064 \frac{1/6}{\Delta a} + 4.8 \frac{1/6}{625} \leq 1,$$

and it is sufficient to choose  $\Delta a > 0.256$ . We adopt

$$\Delta a = \frac{1}{3} \quad (36)$$

The index will be  $i = 0, 1, 2, 3$   $I = 3$ .

### 5.3 Induction formula for $u_1$

$$\begin{aligned} u_1(k+1, i, j) = & u_1(k, i, j) - v_1 \frac{\Delta t}{\Delta a} (u_1(k, i, j) - u_1(k, i-1, j)) + \\ & + D \frac{\Delta t}{(\Delta x)^2} (u_1(k, i, j+1) - 2u_1(k, i, j) + u_1(k, i, j-1)). \end{aligned} \quad (37)$$

Proceeding along the same lines as in case of the model dynamics between E2 and E4, we obtain the formula

$$\begin{aligned} U_1(k+1, j) = & U_1(k, j) - v_1 \frac{\Delta t}{\Delta a} \{u_1(k, I, j) - u_1(k, -1, j)\} + \\ & + D \frac{\Delta t}{(\Delta x)^2} \{U_1(k, j+1) - 2U_1(k, j) + U_1(k, j-1)\}, \end{aligned} \quad (38)$$

where  $U_1(l, r) = \sum_{i=0}^I u_1(l, i, r)$  is the total number of interphasic cells  $G_1 + S + G_2$  at instant  $l$  and at section  $r$ ,  
 $u_1(k, I, j)$  is the number of cells at the end of the interphase  $G_1 + S + G_2$  at instant  $k$ , and at section  $j$ ,  
 $u_1(k, 0, j)$  is the number of cells starting the interphase  $G_1 + S + G_2$  at instant  $k$ , and at section  $j$ .

As we did it for the model between  $E2$  and  $E4$ , we suppose that there exists a constant  $\alpha(k, j)$  such that

$$u_1(k, 0, j) - u_1(k, I, j) = \alpha(k, j)U_1(k, j), \quad (39)$$

and that between  $E4$  and  $E6$   $\alpha$  is also independent of  $k$ , i.e.  $\alpha(k, j) = \alpha(j)$ .

Replacing in (38), and bearing in mind (39), the values of  $v_1, \Delta t, \Delta x, \Delta a, D$  in (35) and (36), we obtain

$$U_1(k+1, j) = (1 + 0.765\alpha(j))U_1(k, j) + 0.00064 \left( \frac{U_1(k, j+1) + U_1(k, j-1)}{2} - U_1(k, j) \right). \quad (40)$$

## 5.4 Approximation of the induction formula

The term

$$0.00064 \left( \frac{U_1(k, j+1) + U_1(k, j-1)}{2} - U_1(k, j) \right)$$

effectively acquires very small values which, in a first approximation, will be ignored.

Formula (40) takes the following approximate form

$$U_1(k+1, j) = (1 + 0.765\alpha(j))U_1(k, j). \quad (41)$$

Multiplying expressions (40) for  $k = 0, 1, \dots, 287$  we obtain

$$U_1(48hs, j) = (1 + 0.765\alpha(j))^{288} U_1(0, j). \quad (42)$$

$U_1(48hs, j)$  represents the number of cells generated between  $E4$  and  $E6$  by the cells that at  $E4$  were located at section  $j$ . Given that during this period the model length undergoes a 2.4 fold increase, it is assumed that, at  $E6$  these newly generated cells will expand over 2.4 segments of  $25\mu m$  thickness.

We will indicate with  $s$  the abscissa of transverse sections of the model at  $E6$ . At each abscissa  $s = 1, 2, \dots, 230$  will correspond a segment of  $25\mu m$  thickness.

It is assumed that the cells located at the  $j$ -th segment at  $E4$  give origin to cells that, at  $E6$ , occupy the part of the model corresponding to  $2.4(j-1) \leq s < 2.4j$ ,  $j = 1, 2, \dots, 96$ .

$U_1(48hs, s)$  indicates the number of cells located at the segment of abscissa  $s$  at  $E6$ .

$\bar{U}_1(48hs, s)_{s=2.4(j-1)}^{s=2.4j}$  will indicate the average number of cells at the segment of abscissa  $s$  at  $E6$ , for  $2.4(j-1) \leq s < 2.4j$ . We have

$$\bar{U}_1(48hs, s)_{s=2.4(j-1)}^{s=2.4j} = \frac{U_1(48hs, j)}{2.4} \quad (43)$$

The values of  $j$  will be  $j = 1, 2, \dots, 96$ , which correspond to 96 segments of the model at  $E4$ .



In order to abbreviate notations, in the following paragraphs, the formula

$$\bar{U}_1(48hs, s) = \bar{U}_1(48hs, s)_{s=2.4(j-1)}^{s=2.4j}$$

will simply be written as  $\bar{U}_1(48hs, s)$  for the corresponding value of  $j$ .

We will have

$$\bar{U}_1(48hs, s) = \frac{1}{2.4} (1 + 0.765\alpha(j))^{288} U_1(0, j)$$

for  $2.4(j-1) \leq s < 2.4j$ ,  $j = 1, 2, \dots, 96$  or

$$\bar{U}_1(48hs, s) = \beta(j)U_1(0, j)$$

where

$$\beta(j) = \frac{1}{2.4} (1 + 0.765\alpha(j))^{288} \quad (44)$$

We define the cell proliferation at segment  $j$  as the quotient

$$prol(j) = \frac{U_1(48hs, j)}{U_1(0hs, j)}$$

Since

$$U_1(48hs, j) = 2.4 \bar{U}_1(48hs, s)$$

we obtain

$$prol(j) = 2.4\beta(j).$$

The  $\beta(j)$  coefficient is proportional to the cell proliferation at section  $j$ .

Experimental data indicate that during the interval  $E4 - E6$  cell proliferation is not uniformly distributed along the cephalic-caudal axis. Thus, a coefficient  $\beta(j)$  is introduced by means of the following simplified expression

$$\beta(j) = \begin{cases} 0, 1j + 0.525 & \text{if } j = 1, 2, 3, 4, 5, 6 \\ 1.125 & \text{if } j = 6, 7, \dots, 48 \\ -0.01j + 1.605 & \text{if } j = 48, \dots, 95, 96 \end{cases} \quad (45)$$

From (44) we obtain

$$\alpha(j) = \frac{1}{0.765} \left( (2.4\beta(j))^{\frac{1}{288}} - 1 \right)$$

and replacing by (45) we obtain the final expression for  $\alpha(j)$ .

Finally, we will have

$$\bar{U}_1(48hs, s) = \beta(j)U_1(0, j) \quad \text{for } j = 1, 2, \dots, 96, \quad (46)$$

where  $\beta(j)$  is given by (45).

## 5.5 Number of cells given by formula (43) and comparison with experimental data

Number of cells per section  $s$  is reported in TABLE 4 (Annexe)

## 5.6 Approximation by linear interpolation of the cell number in intermediate segments of the model at $E6$

We will denote by  $U_1(48hs, s) = U_1(s)$  the number of cells at segment  $s$  of the model at  $E6$  obtained by linear interpolation between values of TABLE 4.

We will have

$$U_1(s) = \left(1 - \frac{s - 2.4(j-1)}{2.4}\right) \bar{U}_1(2.4(j-1)) + \frac{s - 2.4(j-1)}{2.4} \bar{U}_1(2.4j) \quad (47)$$

for sections of the model in  $E6$ , with abscissæ,  $2.4(j-1) \leq s < 2.4j$ .

We apply formula (47) to calculate  $U_1(s)$  for integer values of  $s$ .

Total number of cells per section  $s$  in  $E4$  is reported in TABLE 5 (Annexe) and Fig. 2.

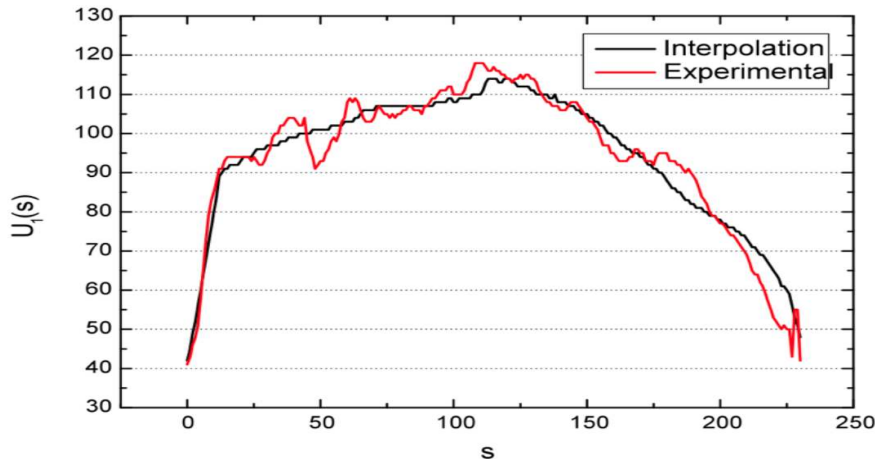


Figure 2: Total number of cells per section  $s$  in  $E6$ .

## 6 Discussion

### 6.1 How to analyze the developmental role of cell proliferation in a developing system?

The proliferation of NE cells is not only involved in generating the appropriate number of cells in the CNS. A spatially organized cell proliferation activity also possesses morpho- and histogenetic effects [32, 23, 33, 34]. These developmental effects should depend on the temporal dynamics and the spatial organization of the proliferative activity. Taking this notion in mind, the present paper attempts at designing a mathematical PDE-based model with the ability to describe the increase in cell number as a function of both time and space. The model emphasizes the occurrence of (a) changes in the cell cycle duration as function of time and (b) changes in the density of NE cells as a function of time and space.

In our opinion this sort of PDE-based models can be useful tools to clearly understand the cell proliferation spatial and temporal organization and to design formal models of cell proliferation that could account for its developmental effects.

This paper is a first step towards designing a model that could help to globally describe the temporal and spatial organization of the cell proliferation and to understand

its specific roles in the genesis of CNS patterning and its supracellular complexity.

A current notion in theoretical biology is that the supracellular complexity is the net result of several DCBs operating simultaneously and interactively [2]. The corticogenesis requires the integrated operation of several DCBs. Such organized operation requires spatially organized cell signaling processes which do not only control cell proliferation and migration but also establish cell-type differences and also determines tissue patterning [35]. It is currently accepted that these kinds of directive influences are mediated by gradients of specific signaling molecules, the so-called morphogens. A developmental gradient may be defined as an asymmetry in morphogen concentration with a maximum at the site of the morphogen-secreting cells and decaying as a function of the distance from that site [3].

As an example, the non-uniform distribution of a morphogen in an embryonic field differentially determines the fate and phenotype of those cells [37]. In a same way, it can be proposed that the non-uniform spatial distribution of developmentally active signals influencing the proliferative activity should result in a non-uniform distribution of the density of proliferating cells and also in the existence of space-dependent differences in cell density along a defined spatial axis.

## 6.2 Stochastic approaches on the NE cell proliferation dynamics during the OT corticogenesis

In the absence of directive influences a DCB should operate at random. In such condition a biological time or space signal representing the DCB operation or representing a quantifiable direct developmental effect should display a random white noise-like dynamics.

A previous work devoted to characterize the postmitotic neuronal migration dynamics in the developing OT by means of standardized methods of frequency signal analyses [38], proposes a mathematical expression that globally describes how cell migration operates in space. This expression includes deterministic and stochastic components. The deterministic component can be described as the sum of a linear global trend and a Fourier series while the stochastic component can be described as a non-correlated process (white noise).

It is reasonable considering that, in the absence of directive influences the NE cells proliferation should operate at random and, consequently, the intercalation of newly generated cells into the whole system should be uniformly distributed. Under such condition the OT growth should proceed as a uniformly distributed tangential expansion; a characteristic that does not occur during the normal OT development. In a previous paper, aiming at determining whether the NE cell proliferation exhibits a spatial organization, spatial records of proliferating (mitotic) NE cells were analyzed within the framework provided by the stochastic point process model [16]. These analyses showed that the NE cell proliferation records can also be described as comprising two components: (a) a non-stationary one that can be deterministically described as a trend plus (b) a stationary non-correlated stochastic point process. The deterministic trend can be interpreted as manifestation of an asymmetrically distributed controlling influence that installs space-dependent differences in the NE cells proliferation rate along the cephalic-caudal axis.

A more recent paper [39] indicates that the dynamics of a detrended NE cell proliferation record does not accurately coincide with a non-correlated stochastic point process with uniform probability distribution of intermitotic intervals. By contrast, stochastic point processes simulated with intermitotic intervals displaying lognormal, exponential and gamma distribution better approximate the NE cell proliferation

records than those simulated with a uniform probability distribution.

### 6.3 The present conclusions on the temporal and spatial organization of the NE cells proliferation during the OT corticogenesis. Morphogenetic correlations

The PDE-based model presented in this paper was specifically design to obtain a better and more realistic characterization of the deterministic behavior exhibited by the proliferating NE cells. In our opinion, the present results appropriately describe the spatial organization of the proliferation activity and explain the differential growth the developing OT displays between *E2E6*. The model accurately estimates the increase in the total number of cells along the defined temporal windows (*E2-E4* and *E4-E6*). These increases closely coincide with values of cell density empirically estimated by means of cells counts performed on histological sections of developing OTs.

With regards to the spatial organization of the NE cells proliferation, the model appropriately reproduces the empirically recorded position-dependent differences in cell density observed in successively ordered  $25\mu m$  length windows along the OT cephalic-caudal axis. In fact, figures 1 and 2 show that the cell density (number of cells / each  $25\mu m$  windows) significantly changes as a function of their position along the cephalic-caudal axis. A comparison between these figures shows that, considering the OT as a whole, the NE cells proliferation significantly decreases as a function of the time (compare the profile of cell density values obtained at the end of the interval *E2-E4* with the profile of cell density at the end of the interval *E4-E6*).

It is interesting remarking that, apart from reproducing the increase in the total number of cells, the profiles of cell density as a function of the space also allows accounting for some morphogenetic events. In fact, between *E2E6* the OT undergoes a remarkable change in shape and size. The differences in cell proliferation as a function of their position along the cephalic-caudal axis should result in a differential intercalation of new cells within the OT wall along that spatial axis. The addition of fewer new cells at both extremes of the OT longitudinal axis should imply a lower OT growth rate at those sites. It must be noted that the low rate of NE cell proliferation at both ends of the cephalic-caudal axis reveals a low NE proliferation rate along the dorsal midline. This lower proliferation activity should also produce a lower growth rate along the entire dorsal midline. This assumption is corroborated by the fact that between *E2E6* a deep medial groove appears along the dorsal midline. This change is a relevant morphogenetic effect since the appearance of a medial groove leads to the constitution of the right and the left OT hemispheres.

All these data about (a) the changes in the number of cells as a function of the time, (b) the space-dependent differences in NE cell proliferation along the cephalic-caudal axis together with (c) the adequacy between the changes in size and shape that can be inferred from the model results, strongly suggest that the PDE-based model here presented can be appropriately and advantageously used to more precisely describe and to more deeply understand the developmental effects of a spatially and temporally organized NE cell proliferation during the OT corticogenesis.

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# ANNEXE

TABLE 1

Section	Number intermitotic cells	Number of mitotic cells
$j = 1$	35	2
2	42	1
3	44	1
4	46	1
5	47	1
6	49	1
7	49	1
8	50	2
9	52	1
10	53	1
11	54	1
12	54	1
13	53	1
14	51	1
15	51	1
16	49	1

TABLE 2

<i>Section</i>	$U_1(0, j)$	$\bar{U}_1(48hs, s)$ <i>according (30)</i>	$\bar{U}_1(48hs, s)$ <i>experimental data</i>
$j = 1$	35	68	78
2	42	81	86
3	44	85	86
4	46	89	87
5	47	91	91
6	49	95	94
7	49	95	98
8	50	96	100
9	52	100	101
10	53	102	101
11	54	104	102
12	54	104	103
13	53	102	101
14	51	98	101
15	51	98	101
16	49	95	72



TABLE 3

<i>Section order</i>	<i>N<sup>er</sup>cells of the Model at E4 obtained by interpolation</i>	<i>N<sup>er</sup>cells of the Tectum at E4 according to experimental data</i>
$s = 1$	68	65
2	70	72
3	72	79
4	75	82
5	77	84
6	79	86
7	81	88
8	82	88
9	82	86
10	84	86
11	84	85
12	85	85
13	85	85
14	86	85
15	86	85
16	87	85
17	88	86
18	88	87
19	89	87
20	89	88
21	90	88
22	90	87
23	90	87
24	91	89
25	91	89
26	92	90
27	92	91
28	93	91
29	94	93
30	94	93
31	95	95
32	95	95
33	95	95
34	95	93
35	95	93
36	95	96
37	95	96
38	95	97
39	95	98
40	96	98
41	96	99
42	97	99
43	96	100
44	97	99
45	97	100
46	98	99
47	99	99
48	99	102
49	100	101

$s = 50$	100	102
51	101	101
52	101	101
53	101	102
54	102	101
55	102	102
56	102	101
57	103	102
58	103	101
59	103	102
60	104	102
61	104	102
62	104	102
63	104	102
64	104	102
65	104	103
66	104	102
67	104	103
68	104	102
69	103	103
70	103	103
71	103	103
72	102	102
73	102	102
74	101	101
75	101	101
76	100	100
77	99	100
78	99	100
79	98	99
80	98	100
81	98	102
82	98	102
83	98	103
84	98	103
85	98	104
86	98	103
87	97	103
88	97	99
89	96	94
90	95	90
91	95	83
92	93	77
93	90	69
94	88	73
95	86	67
96	83	60

TABLE 4

		$E4$ $U_1(0, j)$	$E6$ $\bar{U}_1(48hs, s)_{s=2.4(j-1)}^{s=2.4j}$ <i>according to (43)</i>	$E6$ $\bar{U}_1(48hs, s)_{s=2.4(j-1)}^{s=2.4j}$ <i>experimental data</i>
$j = 1$	$0 \leq s < 2.4$	68	43	43
2	$2.4 \leq s < 4.8$	70	51	48
3	$4.8 \leq s < 7.2$	72	59	62
4	$7.2 \leq s < 9.6$	75	69	78
5	$9.6 \leq s < 12$	77	79	86
6	$12 \leq s < 14.4$	79	89	91
7	$14.4 \leq s < 16.8$	81	91	94
8	$2.4 \leq s < 4.8$	82	92	94
9	$4.8 \leq s < 7.2$	82	92	94
10	$7.2 \leq s < 9.6$	84	94	94
11	$9.6 \leq s < 12$	84	94	93
12	$12 \leq s < 14.4$	85	96	93
13	$28.8 \leq s < 31.2$	85	96	94
14	$31.2 \leq s < 33.6$	86	97	98
15	$33.6 \leq s < 36$	86	97	101
16	$36 \leq s < 38.4$	87	98	103
17	$38.4 \leq s < 40.8$	88	99	104
18	$40.8 \leq s < 43.2$	88	99	102
19	$43.2 \leq s < 45.6$	89	100	100
20	$45.6 \leq s < 48$	89	100	95
21	$48 \leq s < 50.4$	90	101	91
22	$50.4 \leq s < 52.8$	90	101	94
23	$52.8 \leq s < 55.2$	90	101	97
24	$55.2 \leq s < 57.6$	91	102	99
25	$57.6 \leq s < 60$	91	102	103
26	$60 \leq s < 62.4$	92	104	108
27	$62.4 \leq s < 64.8$	92	104	108
28	$64.8 \leq s < 67.2$	93	105	105
29	$67.2 \leq s < 69.6$	94	106	103
30	$69.6 \leq s < 72$	94	106	105
31	$72 \leq s < 74.4$	95	107	107
32	$74.4 \leq s < 76.8$	82	92	94
33	$76.8 \leq s < 79.2$	82	92	94
34	$79.2 \leq s < 81.6$	84	94	94
35	$81.6 \leq s < 84$	84	94	93
36	$84 \leq s < 86.4$	85	96	93
37	$86.4 \leq s < 88.8$	95	107	106
38	$88.8 \leq s < 91.2$	95	107	107
39	$91.2 \leq s < 93.6$	95	107	109
40	$93.6 \leq s < 96$	96	108	110
41	$96 \leq s < 98.4$	96	108	111
42	$98.4 \leq s < 100.8$	97	109	111
43	$100.8 \leq s < 103.2$	96	108	110
44	$103.2 \leq s < 105.6$	97	109	111
45	$105.6 \leq s < 108$	97	109	115
46	$108 \leq s < 110.4$	98	110	118
47	$110.4 \leq s < 112.8$	98	110	118
48	$112.8 \leq s < 115.2$	99	111	116

$j=49$	$115.2 \leq s < 117.6$	100	108	110
50	$117.6 \leq s < 120$	100	109	111
51	$120 \leq s < 122.4$	101	109	115
52	$122.4 \leq s < 124.8$	101	110	118
53	$124.8 \leq s < 127.2$	101	110	118
54	$127.2 \leq s < 129.6$	102	111	116
55	$129.6 \leq s < 132$	102	108	113
56	$132 \leq s < 134.4$	102	107	110
57	$134.4 \leq s < 136.8$	103	107	108
58	$136.8 \leq s < 139.2$	103	106	107
59	$139.2 \leq s < 141.6$	103	105	106
60	$141.6 \leq s < 144$	104	105	107
61	$144 \leq s < 146.4$	104	102	108
62	$146.4 \leq s < 148.8$	104	102	107
63	$148.8 \leq s < 151.2$	104	101	105
64	$151.2 \leq s < 153.6$	104	100	103
65	$153.6 \leq s < 156$	104	99	100
66	$156 \leq s < 158.4$	104	98	97
67	$158.4 \leq s < 160.8$	104	97	95
68	$160.8 \leq s < 163.2$	104	96	94
69	$163.2 \leq s < 165.6$	103	94	93
70	$165.6 \leq s < 168$	103	93	94
71	$168 \leq s < 170.4$	103	92	96
72	$170.4 \leq s < 172.8$	102	90	95
73	$172.8 \leq s < 175.2$	102	89	93
74	$175.2 \leq s < 177.6$	101	87	94
75	$177.6 \leq s < 180$	101	86	95
76	$180 \leq s < 182.4$	100	86	94
77	$182.4 \leq s < 184.8$	99	83	92
78	$184.8 \leq s < 187.2$	99	82	91
79	$187.2 \leq s < 189.6$	98	80	90
80	$189.6 \leq s < 192$	98	79	89
81	$192 \leq s < 194.4$	98	78	84
82	$194.4 \leq s < 196.8$	98	77	81
83	$196.8 \leq s < 199.2$	98	76	79
84	$199.2 \leq s < 201.6$	98	75	78
85	$201.6 \leq s < 204$	98	74	76
86	$204 \leq s < 206.4$	98	73	74
87	$206.4 \leq s < 208.8$	97	71	72
88	$208.8 \leq s < 211.2$	97	70	69
89	$211.2 \leq s < 213.6$	96	69	66
90	$213.6 \leq s < 216$	95	67	63
91	$216 \leq s < 218.4$	95	66	60
92	$218.4 \leq s < 220.8$	93	64	55
93	$220.8 \leq s < 223.2$	90	61	51
94	$223.2 \leq s < 225.6$	88	58	50
95	$225.6 \leq s < 228$	86	56	47
96	$228 \leq s < 230.4$	83	54	53

TABLE 5

<i>Section</i> $s$	$U_1(s)$ <i>according to (44)</i>	$U_1(s)$ <i>according experiments</i>
0	42	41
1	45	43
2	49	46
3	52	48
4	56	51
5	60	58
6	64	65
7	68	73
8	72	79
9	76	83
10	80	85
11	84	88
12	89	91
13	90	91
14	91	93
15	91	94
16	92	94
17	92	94
18	92	94
19	92	94
20	93	94
21	94	94
22	94	94
23	94	94
24	94	93
25	95	94
26	96	93
27	96	92
28	96	92
29	96	93
30	97	95
31	97	96
32	97	99
33	97	100
34	97	100
35	98	102
36	98	102
37	98	103
38	99	104
39	99	104
40	99	104
41	99	103
42	100	102
43	100	102
44	100	104
45	100	98
46	100	96
47	101	94
48	101	91
49	101	92
50	101	93
51	101	93

<i>Section</i>	$s$	$U_1(s)$ <i>according to (44)</i>	$U_1(s)$ <i>according experiments</i>
52		101	95
53		101	96
54		102	98
55		102	99
56		102	98
57		102	100
58		102	103
59		103	104
60		103	108
61		103	109
62		103	108
63		104	109
64		104	108
65		105	106
66		106	104
67		106	103
68		106	103
69		106	103
70		106	104
71		107	106
72		107	107
73		107	106
74		107	105
75		107	105
76		107	104
77		107	105
78		107	104
79		107	105
80		107	105
81		107	106
82		107	106
83		107	107
84		107	107
85		107	106
86		107	106
87		107	106
88		107	105
89		107	107
90		107	107
91		107	108
92		107	109
93		108	109
94		108	110
95		108	111
96		108	111
97		108	111
98		109	112
99		109	112
100		108	110
101		108	110
102		109	110
103		109	110
104		109	111
105		109	113

<i>Section</i>	$s$	$U_1(s)$ <i>according to (44)</i>	$U_1(s)$ <i>according experiments</i>
106		109	114
107		110	116
108		110	118
109		110	118
110		110	118
111		111	118
112		113	117
113		114	116
114		114	116
115		114	117
116		114	116
117		113	116
118		113	115
119		114	115
120		114	114
121		114	114
122		113	113
123		113	114
124		112	114
125		112	115
126		112	114
127		112	115
128		112	115
129		111	114
130		111	114
131		110	113
132		110	111
133		110	110
134		110	109
135		110	108
136		109	108
137		109	107
138		110	107
139		108	107
140		108	106
141		108	106
142		108	106
143		107	107
144		107	108
145		107	108
146		106	108
147		106	107
148		105	106
149		105	105
150		105	104
151		104	103
152		104	103
153		103	102
154		103	101
155		102	99
156		102	97
157		101	97
158		100	97
159		100	95
160		99	95

<i>Section</i>	$s$	$U_1(s)$ <i>according to (44)</i>	$U_1(s)$ <i>according experiments</i>
161		99	94
162		98	93
163		97	93
164		97	93
165		96	93
166		96	94
167		95	94
168		95	96
169		95	96
170		94	95
171		94	95
172		93	93
173		92	93
174		92	93
175		91	92
176		91	94
177		90	95
178		90	95
179		89	95
180		88	95
181		87	93
182		86	93
183		86	93
184		85	92
185		85	92
186		84	91
187		83	90
188		83	91
189		82	90
190		82	89
191		81	88
192		81	86
193		81	84
194		80	83
195		80	82
196		79	80
197		79	79
198		79	79
199		78	78
200		78	77
201		77	77
202		77	76
203		76	75
204		76	74
205		76	74
206		75	73
207		75	72
208		74	71
209		74	70
210		73	69



<i>Section</i>	$s$	$U_1(s)$ <i>according to (44)</i>	$U_1(s)$ <i>according experiments</i>
211		72	67
212		71	65
213		71	64
214		70	64
215		69	62
216		69	61
217		68	59
218		67	57
219		66	55
220		65	53
221		64	52
222		63	51
223		61	50
224		61	51
225		60	50
226		59	50
227		56	43
228		53	55
229		51	55
230		48	42